

87261

S/033/60/037/006/020/022  
E032/E514

An Interference Polarization Filter for Astrophysical Studies of  
the Sun in the K-Line of Ionized Calcium

photographs were obtained in the  $KCa^+$  line using the horizontal  
solar telescope (diameter of image of the Sun 16 cm). Typical  
photographs obtained are given. Acknowledgments are expressed to  
Academician V. P. Linnik for interest in this work. There are  
7 figures and 10 references: 8 Soviet, 2 non-Soviet.

ASSOCIATION: Gosudarstvennyy opticheskiy institut imeni  
S. I. Vavilova (State Optical Institute imeni  
S. I. Vavilov)

SUBMITTED: May 4, 1960

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87261

S/033/60/037/006/020/022

E032/E514

An Interference Polarization Filter for Astrophysical Studies of  
the Sun in the K-Line of Ionized Calcium

Fig.1

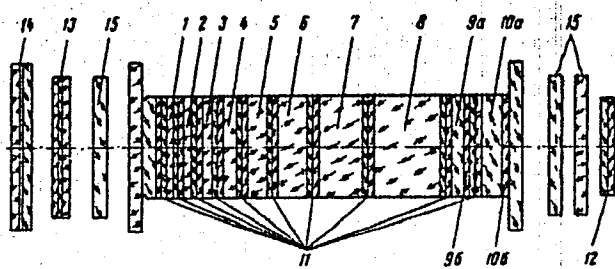


Рис. 1

Card 4/4

IOFFE, S.B.; SMIRNOVA, T.A.

Achromatic wave plates. Opt. i spektr. 16 no. 5 894-897 My '64.  
(MIRA 17:9)

L 25528-66 EWP(e)/EWT(m)/EWP(b) WH

ACC NR: AP5025861

SOURCE CODE: UR/0020/65/164/004/0793/0795

AUTHOR: Ioffe, S. B.; Drichko, N. M.

ORG: none

TITLE: Phase dispersive interference-polarization filters

SOURCE: AN SSSR. Doklady, v. 164, no. 4, 1965, 793-795

TOPIC TAGS: optic filter, polarizing filter

ABSTRACT: The properties of interference-polarization filters depend basically on the magnitude of the double refraction index and the thickness of the layers, the dispersion of the double refraction index being of secondary importance. In the present article the authors describe a new phase dispersive interference-polarization filter in which the fundamental effect is due to dispersion properties of the materials. Use of different crystalline materials exhibiting varying degrees of dispersion permits the production of passband spectral domains, shown in Figures 1 and 2, with completely novel characteristics.

Card 1/4

UDC: 681.40

L 15528-66

ACC NR: AP5025861

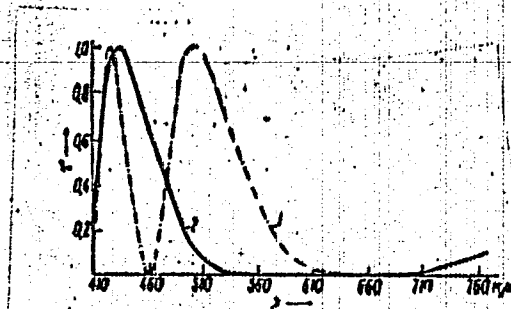


Fig. 1 Passband curves of a two-component phase dispersive interference-polarization filter 1 -  $\lambda_0 = 656.3 \text{ m}\mu$ ,  $N_{01} =$  (magnitude of the wave length difference appearing within the 1st plate for a wavelength  $\lambda_0$ ) = 79.11,  $N_{02} = 77.61$ ,  $n_1 = n_2 = 0$ ,  $v_1 = 45^\circ$ ,  $v_2 = -45^\circ$  ( $n_1$  and  $n_2$  - orientation angles of the polarizer and analyzer;  $v_1$  and  $v_2$  - orientation angles of the plates); 2 -  $\lambda_0 = 656.3 \text{ m}\mu$ ,  $N_{01} = 53.24$ ,  $N_{02} = 51.74$  for the same orientations. Index 1 refers to quartz plates, 2 to spat plates.

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L 15528-66

ACC NR: AP5025861

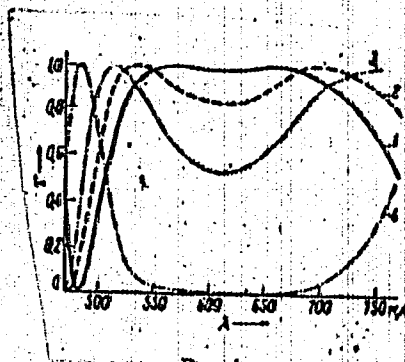


Fig. 2 Passband curves of a two-component phase dispersive interference-polarization filter with an electrooptical phase changing device. 1 -  $\phi_{m+1} = 0$ ; 2 -  $0.2\pi$ ; 3 -  $0.4\pi$ ; 4 -  $\phi_{m+1} = \pi$  ( $\phi_{m+1}$  - phase difference appearing in the  $K_{m+1}$  plate.)  $\lambda_0 = 656.3 \text{ m}\mu$ ,  $N_{01} = 134.35$ ,  $N_{02} = 129.35$ ,  $n_1 = n_2 = 0$ ,  $v_1 = +45^\circ$ ,  $v_2 = -45^\circ$ .

Card 3/4

L 15528-66

ACC NR: AP5025861

The paper was presented by Academician V. P. Linnik, 5 Feb. 65. Orig. art. has: 6 formulas and 4 figures. J

SUB CODE: 20 / SUBM DATE: 11Feb65 / ORIG REF: 001 / OTH REF: 002

  
Card 4/4

IOFFE, Sh.I.

The AChV-Sh carding and knitting unit. Biul. tekhn.-ekon. inform.  
no.10:55-57 '59. (MIRA 13:3)  
(Knitting machines)



S/079/62/032/002/001/011  
D204/D303

AUTHORS: Chernyshev, Ye., A, Tolstikova, N.G., Ioffe, S.L. and Petrov, A.D.

TITLE: Interaction of disilanes with chlorobenzene in the vapor phase

PERIODICAL: Zhurnal obshchey khimii, v. 32, no. 2, 1962, 369-374

TEXT: A continuation of earlier work concerned with the preparation of organochlorosilanes. In the present paper the authors describe the reactions of chlorobenzene with hexamethyl -, pentamethyl chloro-, tetramethyl dichloro- and hexachlorodisilanes. The reactions were studied by passing mixtures of PhCl (2 moles) and the corresponding disilane (1 mole) through a silica tube at 500-600 C. The reagents were in the hot zone for 30-35 secs. The products were then condensed and analyzed. Full experimental details are given. The interaction of PhCl with hexamethyl disilane at 500 and 550 C yielded  $(CH_3)_3SiCl$ ,  $C_6H_6$ ,  $(CH_3)_3Si.Ch_2.Si(CH_3)_2Cl$  ✓

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S/079/02/032/002/001/011  
D204/D303

Interaction of disilanes ...

and  $C_6H_5 \cdot C_6H_5$ . Small quantities of  $Cl(CH_3)_2Si \cdot CH_2 \cdot Si(CH_3)_2Cl$  and  $(CH_3)_3SiC_6H_5$  were also formed at  $600^\circ C$ . The reaction with pentamethyl chloro-disilane gave  $(CH_3)_3SiCl$ ,  $(CH_3)_2SiCl_2$ ,  $C_6H_6$ ,  $Cl(CH_3)_2Si \cdot CH_2 \cdot Si(CH_3)_2Cl$ ,  $Cl(CH_3)_2Si \cdot CH_2 \cdot Si(CH_3)_2Cl_2$ ,  $(CH_3)_3SiC_6H_5$ ,  $Cl(CH_3)_2SiC_6H_5$  and  $C_6H_5 \cdot C_6H_5$ . No thermal rearrangement of pentamethyl chloro-disilane was observed, in contrast to the hexamethyl disilane. The action of  $PhCl$  on tetra methyl dichlorodisilane (at  $600^\circ C$  only) resulted in  $(CH_3)_2SiCl_2$ ,  $C_6H_6$ ,  $Cl(CH_3)_2Si \cdot CH_2 \cdot Si(CH_3)_2Cl_2$ ,  $Cl(CH_3)_2SiC_6H_5$  and  $C_6H_5 \cdot C_6H_5$ . In addition to the above listed compounds, high-boiling residues were formed in the 3 cases. The nature of the products obtained is discussed and it is concluded that these reactions proceed by a free radical chain mechanism. The interactions begin with a displacement of an H-atom in the disilane by a silyl or a phenyl radical, followed by rearrangement of  $\rightarrow Si-Si-CH_2$  into

$\rightarrow Si-CH_2-Si \cdot$ . The interaction, at  $580^\circ C$ , of  $PhCl$  with

Card 2/3

IOFFE, S. L.; TARTAKOVSKIY, V. A.; NOVIKOV, S. S.

Mechanism of the reduction of carbonyl-containing compounds with  
diborane solution in tetrahydrofuran. Izv AN SSSR Ser Khim no. 4:  
622-631 Ap '64. (MIRA 17:5)

1. Institut organicheskoy khimii im. N. D. Zelinskogo AN  
SSSR.

IOFFE, S.L.; TARTAKOVSKIY, V.A.; MEDVEDEVA, A.A.; NOVIKOV, S.S.

Reduction of oximes with diborane solution in tetrahydrofuran.  
Izv. AN SSSR. Ser. khim. no.8:1537-1538 Ag '64.

1. Institut organicheskoy khimii im. N.D. Zelinskogo AN SSSR. (MIRA 17:9)

IOFFE, S.L.; TARTAKOVSKIY, V.A.; NOVIKOV, S.S.

Selective reduction of aliphatic functional nitro compounds.  
Usp. khim. 35 no.1:43-69 Ja '66. (MIRA 19:1)

1. Institut organicheskoy khimii AN SSSR imeni N.D. Zelinskogo.

10

Reactions of nitroso compounds with nitric oxide. A. N. Nesmeyanov and S. T. Ioffe. *J. Gen. Chem.* (U. S. S. R.) 11, 392-401 (1941).—The C-nitroso group of nitrosodialkylanilines, nitrosoaralkylanilines, N,C-dinitrosoaralkylanilines, 1-nitrosoaphthalene, 4-nitrosoantipyrine and nitrosophenol reacts with NO in accordance with the Bamberger scheme (C. A. 13, 2571) with the formation of the nitrate of the aryl or heterocyclic diazonium compound. The formation of the phenyldiazonium nitrate by the reaction of NOCl with the salt of nitrosophenylhydroxylamine lends support to the Bamberger supposition that the reaction occurs through the addn. of 2 mols. of NO at the double bond of the nitroso group followed by the rearrangement. B. Z. Kamich

10

IOFFE, S. T., et al.

Science

Manual on organomagnesium compounds. v. 3. (Sinteticheskie metody v oblasti metalloorganicheskikh soedinenii, no. 2). Moskva, Izd-vo AN SSSR, 1950.

Monthly List of Russian Accessions, Library of Congress, October 1952. UNCLASSIFIED.

191112

USSR/Chemistry - Organometallic  
Compounds

Sep/Oct 51

"Contributions of Russian Chemists to the Development of Magnesium-Organic /Grignard/ Synthesis (To the 50th Anniversary of Grignard's Synthesis)," S. T. Ioffe, Moscow

"Uspekh Khim" Vol XX, No 5, pp 621-656

Points out that the Kazan' school was the 1st to apply Zn organic compds in synthetic work and that out of 5,000 major contributions dealing with the use of Mg compds in synthesis, 700 were published by Russians (cf. S. T. Ioffe, A. N. Nesmeyanov, "Handbook of Magnesium-Organic Compounds,"

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USSR/Chemistry - Organometallic  
Compounds (Contd) Sep/Oct 51

Moscow, 1950). Discusses use of Zn-org compds, synthesis of aliphatic hydrocarbons and their functional derivs, synthesis of acetylene derivs, application of Mg compds in the fld of terpene chemistry, use of Be-org compds in the synthesis of other organoelemental compds (work by A. N. Nesmeyanov, K. A. Kocheshkov, et al; G. Kamay; P. P. Shorygin et al; etc.), industrial Mg syntheses with the aid of Mg-org compds prep'd in the absence of ether (Shorygin's pioneering work on this type of synthesis and its applications in industrial production of synthetic essential oils), anomalies in Mg-org synthesis.

191112



191714

USSR/Chemistry - Organometallic  
Compounds

Sep/Oct 51

"Review of S. T. Ioffe and A. N. Nesmeyanov's  
'Handbook of Magnesium-Organic Compounds,' Volume  
I-III," Ye. T. Evtus

"Uspekhi Khim" Vol XX, No 5, pp 671, 672

Reviews in some detail this work, which is a part  
of the series "Synthetic Methods in the Field of  
Organometallic Compounds" published under the  
general editorial supervision of Acad A. N.  
Nesmeyanov and K. A. Kocheshkov, Corr Mem, Acad  
Sci USSR. States that all references to

USSR/Chemistry - Organometallic  
Compounds (Contd) Sep/Oct 51

reactions involving Mg-org compds and listed in  
"Chem Zentralblatt," 1899 - 1940, and "Chem Ab-  
stracts," 1941 - Jan 1, 1948, as well as Russian  
and USSR work in this fld not listed by these 2  
journals (altogether more than 13,000 reactions)  
Evdus, the handbook is well published and indexed.  
Published by Press Acad Sci USSR, M-L, 1950.

191714

IOFFE, S. T.

Chemical Abst.  
Vol. 48 No. 4  
Feb. 25, 1954  
General and Physical Chemistry

Theory of tautomeric equilibrium. Effect of solvent on the acidity of the tautomeric forms and position of keto-enol equilibrium. M. I. Kabachnik and S. T. Ioffe (Inst. Org. Chem., Acad. Sci. U.S.S.R., Moscow, U.S.S.R. *Nash S.S.S.R.* 91: 833-6 (1953)). It was previously shown (C.A. 46, 8490f) that the const. of tautomeric equil.  $K_{TS}$  can be expressed as  $K_{TS} = K_{KS}/K_{SS}$ , where  $K_{KS}$  is the const. of acidic ionization of ketonic forms in a given solvent, and  $K_{SS}$  is the const. of ionization of the enolic forms. By the use of the Brønsted concept of protolytic equilibria and soln. of the 2 simultaneous equations:  $pK_{KS} = pK_{KS} + \text{const.}$  and  $pK_{SS} = pK_{SS} + \text{const.}$  gives  $pK_{TS} = pK_{TS} + \text{const.}$  for enol-keto systems in 2 solvents  $S_1$  and  $S_2$ . Exptl. detn. of ionization consts. of tautomeric compds., however, yields only some summary const. of the equil. mixt. (*ibid.* 83, 850 (1953)). However, exptl. detn. of the ionization const. of the equil. system  $K_{KS}$  and detn. of the const. of tautomeric equil.  $K_{TS}$  permits the calcn. of the individual ionization consts. from:  $K_{KS} = K_{KS} / (K_{TS} + 1)$  and  $K_{SS} = K_{KS} / (K_{TS} + 1) / K_{TS}$ . The consts.  $K_{KS}$  were detd. by measurement of pH of partially neutralized solns. by means of a glass electrode. Calcns. of  $K_{SS}$  were made by the formula:  $K_{SS} = ([Na^+] + [H^+] - (K_a/[H^+])) / ([M] - ([Na^+] + [H^+] - (K_a/[H^+])))$ . Plots of  $pK_{TS}$  and  $pK_{KS}$  give linear curves, justifying the above formulation and permitting the calcn. of the individual consts. in solvent pairs of  $H_2O$  and 70% aq. dioxane. The following values are found for the enol content in water and aq. dioxane: acetylacetone 19% and 53%; Et acetoacetate 0.5 and 3%; benzoylacetone 48% and 81%; formylcyclopentanone 40% and 76%; 2-methylindanedione 1% and 6%; Et benzoylacetate 1% and 6%. The calcd. values of  $K_{TS}$ ,  $K_{KS}$ ,  $K_{SS}$ , and  $K_{KS}$  are tabulated. G. M. Kosolapoff.

7-28-54

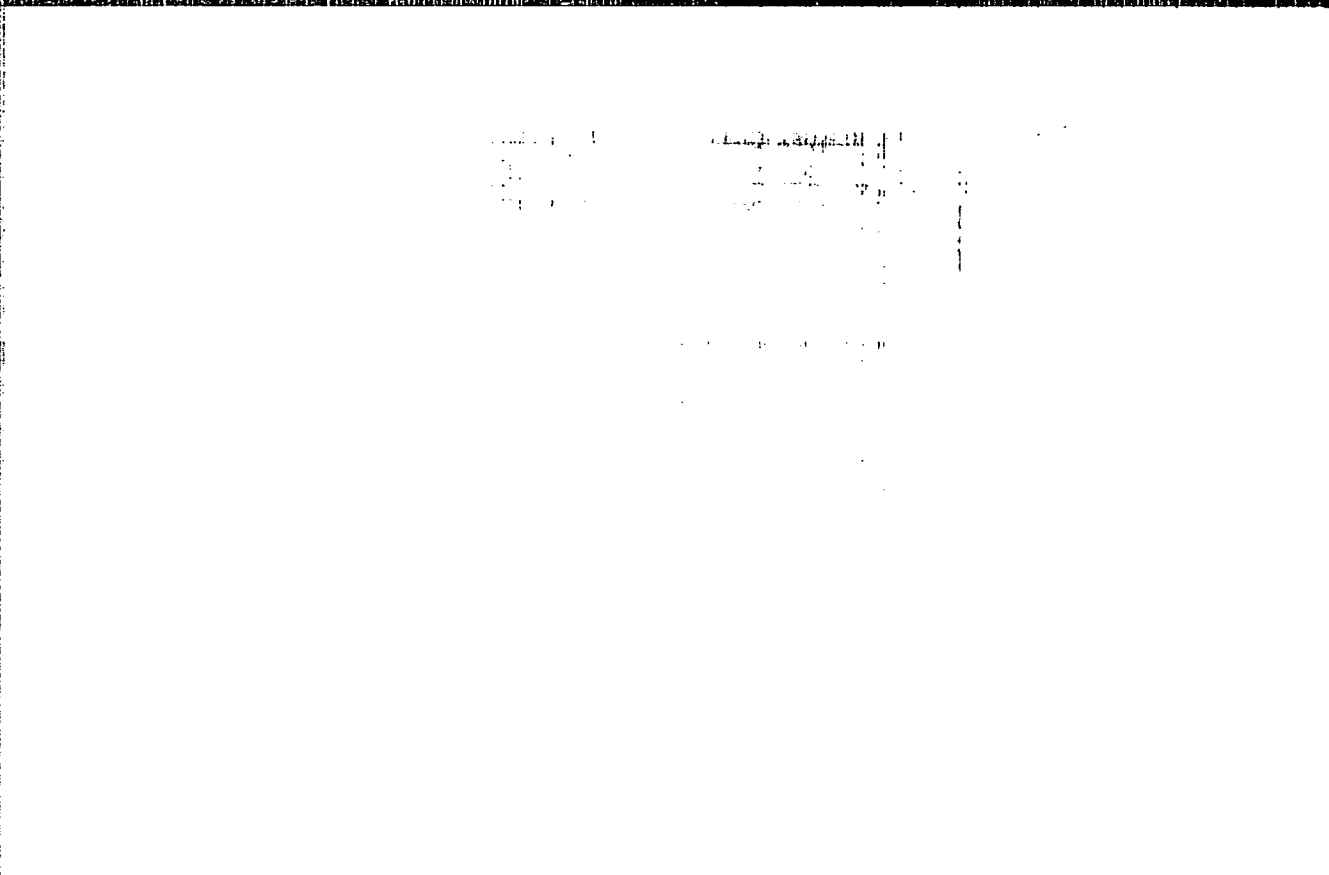
KABACHNIK, M.I.; IOFFE, S.T.; MASTRYUKOVA, T.A.

Theory of tautomeric equilibrium in solutions. Tautomerism of di-  
alkylthiophosphates. Zhur.ob.khim. 25 no.4:684-693 Ap'55.  
(MIRA 8:7)

1. Institut elementoorganicheskikh soedineniy Akademii nauk SSSR.  
(Thiophosphates) (Tautomerism)

"APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000618630003-4

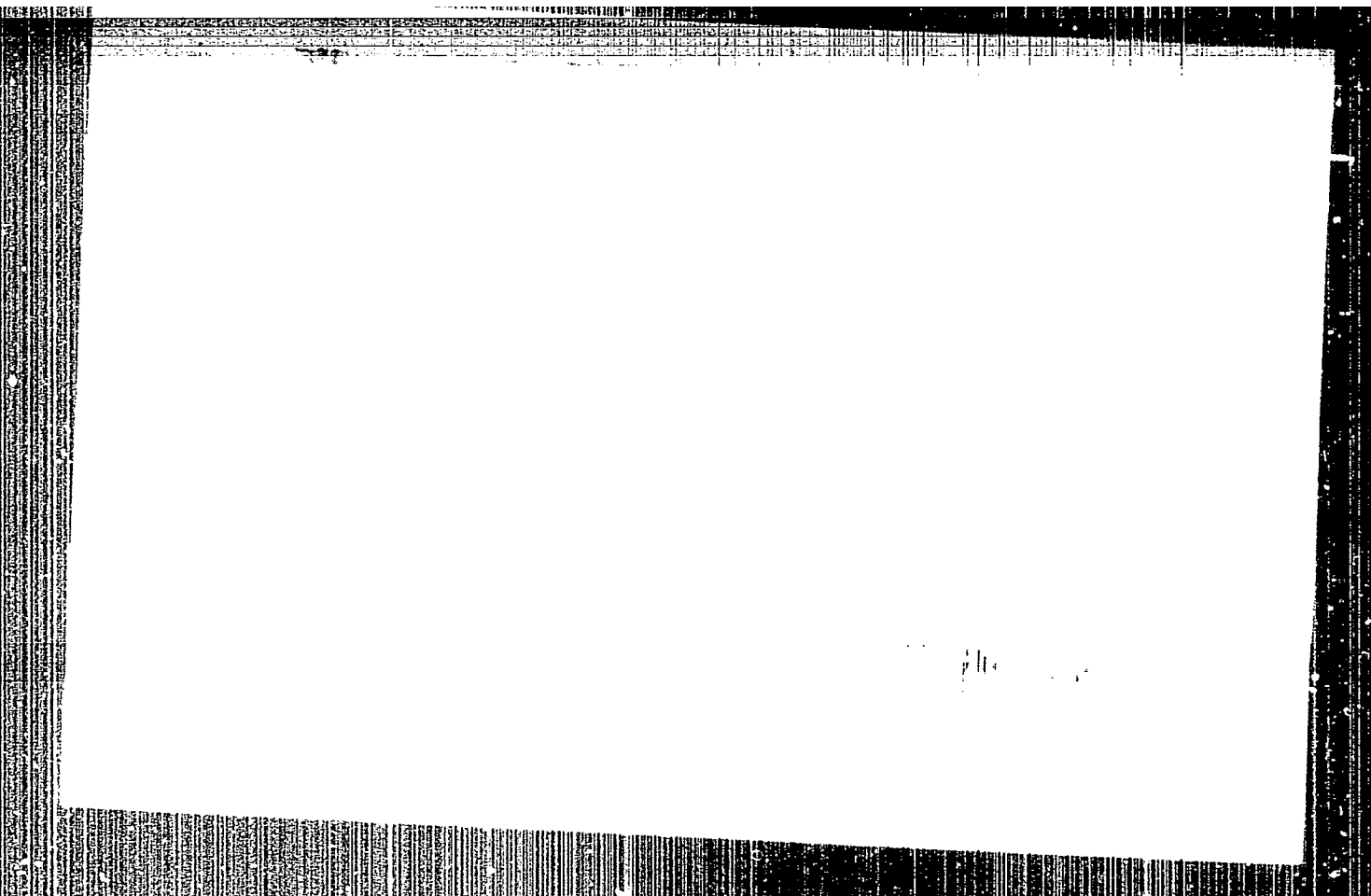


APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000618630003-4"

**"APPROVED FOR RELEASE: 08/10/2001**

**CIA-RDP86-00513R000618630003-4**



**APPROVED FOR RELEASE: 08/10/2001**

**CIA-RDP86-00513R000618630003-4"**

IOFFE, S. T.

"A Study of Tautomerism of Organophosphorous Compounds  
by Potentiometry"  
paper presented at Nn First Conference of Phosphorous Compounds, Kazan,  
8-10 Dec 56

SO: B-3,084, 841

**"APPROVED FOR RELEASE: 08/10/2001**

**CIA-RDP86-00513R000618630003-4**

**APPROVED FOR RELEASE: 08/10/2001**

**CIA-RDP86-00513R000618630003-4"**

IOFFE, S. T. (Institute of Elementary Organic Compounds AS USSR, Moscow)

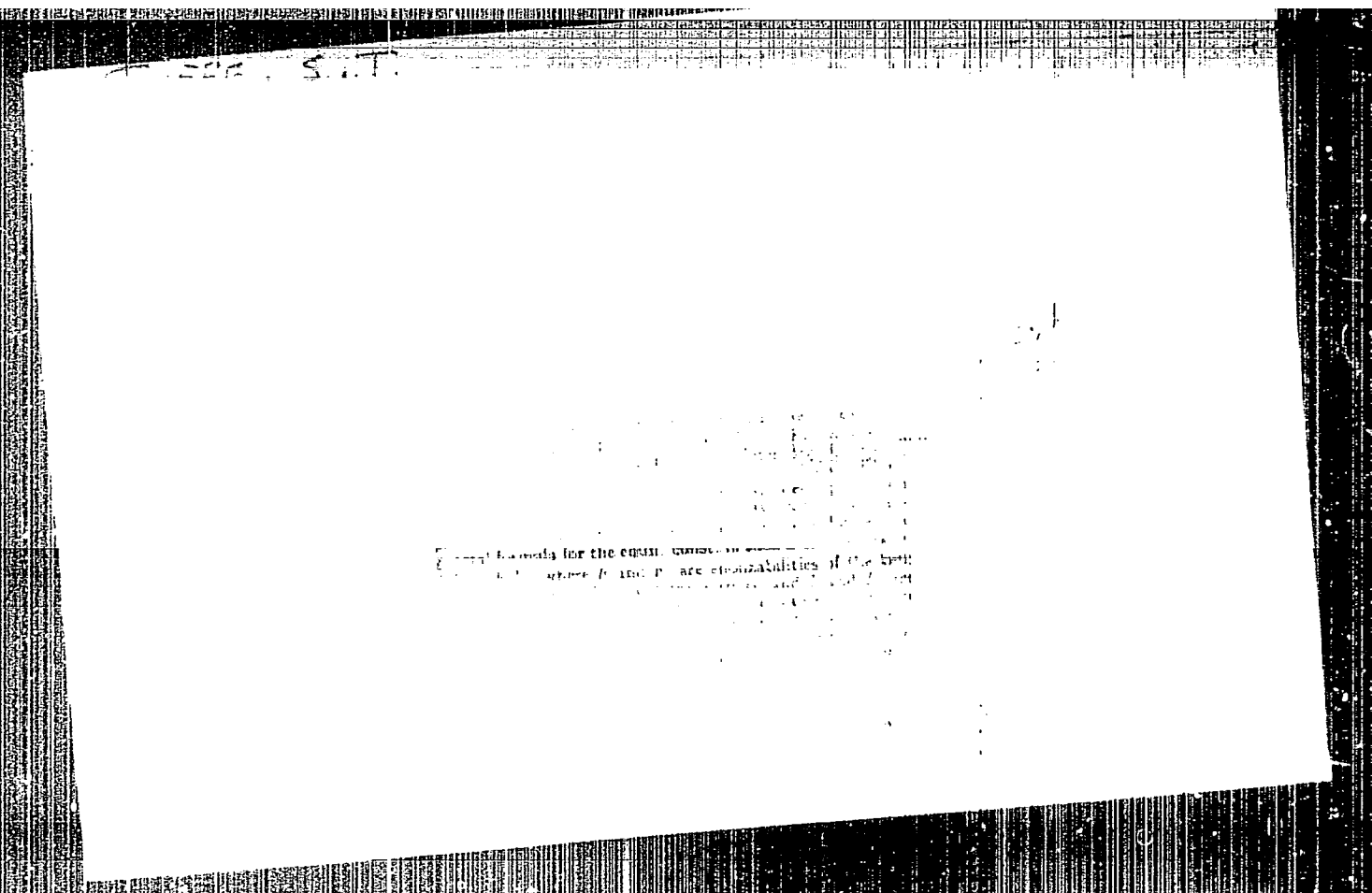
"Research on Tautomerism of Organophosphorus Compounds by the Potentiometric Method" (Issledovaniye tautomerii fosfororganicheskikh soyedineniy potentsiometricheskim metodom)

Chemistry and Uses of Organophosphorous Compounds  
(Khimiya i primeneniye fosfororganicheskikh soyedneniy),  
Trudy of First Conference, 8-10 December 1955, Kazan,  
PP. Published by Kazan Affil. AS USSR, 1957

76-90

Report discussed by A. Ye. Arbuzov, Kazan Aff. AS USSR





*Ioffe, S.T.*

KABACHNIK, M.I.; IOFFE, S.T.; VATSURO, K.V.

Cis-trans-enol tautomerism. Ukr. khim. zhur. 23 no.5:602-614 '57.  
(MIRA 10:11)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.  
(Tautomerism)

62-58-5-16/27

AUTHORS: Kabachnik, M. I., Ioffe, S. T.

TITLE: On the Investigation of Tautomerism in Aprotic Solvents  
(K izucheniye tautomerii v aprotnykh sredakh)

PERIODICAL: Izvestiya Akademii Nauk SSSR Otdeleniye Khimicheskikh Nauk,  
1958, Nr 5, pp. 628 - 630 (USSR)

ABSTRACT: According to Gammet (Reference 1) a difference must be made between ionization and dissociation in solutions. According to Izmaylov (Reference 2) the process of acid-dissociation consists of the following equilibrium-reactions: The solvation of the neutral acid-molecule, the dissociation of the solvate with the formation of solvated ions and the association of these ions in ion-pairs. Already earlier a general potentiometric method for the determination of the constants of the tautomeric equilibrium in conducting media was worked out. In the present report they describe the colorimetric method elaborated by them for the determination of the constants of protolysis in aprotic solvents with low dielectricity-constant. The neutralization-reaction of the acid in aprotic solvents with low dielectric constant leads to the formation of non-dissociated

Card 1/2

On the Investigation of Tautomerism in Aprotic Solvents 62-58-5-16/27

ion-pairs. The values  $n$  are practically equal to 1. The consideration of Brensted's dependence makes the application of the constants of the protolysis for the determination of the constants of the tautomeric equilibrium possible. There are 2 figures, 1 table and 7 references, 3 of which are Soviet.

ASSOCIATION: Institut elementoorganicheskikh soedineniy Akademii nauk SSSR  
( Institute for Elemental-organic Compounds AS USSR)

SUBMITTED: December 17, 1957

1. Organic solvents--Chemical reactions 2. Organic solvents  
--Analysis 3. Acid-base equilibrium--Analysis 4. Colorimetry--Applications

Card 2/2

AUTHORS: Movsesyan, M. Ye., Kabachnik, M. I., SOV/48-22-9-32/40  
Ioffe, S. T., Vatsuro, K. V.

TITLE: Investigation of the Keto-Cis-Trans-Enol Equilibrium by  
Means of Infrared Absorption Spectra (Issledovaniye  
keto-tsis-trans-enol'nogo ravnovesiya pri pomoshchi  
spektrov infrakrasnogo pogloshcheniya)

PERIODICAL: Izvestiya Akademii nauk SSSR, Seriya fizicheskaya, 1958,  
Vol 22, Nr 9, pp 1126 - 1130 (USSR)

ABSTRACT: This paper is an attempt to show the cis-trans enol  
tautomerism in keto-enol compounds by means of infrared  
absorption and to estimate the relative numbers of  
stereoisomeric variants. The investigation covered the  
infrared absorption spectra of acetic ester, of ethyl  
ester, of cyclohexanonic- and of cyclopentanone carboxylic  
acids, of  $\alpha$  secondary butyl acetic ester, and of formyl  
phenyl acetic ester. Chloroform, benzene, toluene,  
diethyl ether, carbon tetrachloride and n-hexane served  
as solvents. The relative intensities of the absorption  
bands of the keto- and enol variants of the substances  
in question were compared with the equilibrium constants

Card 1/3

Investigation of the Keto-Cis-Trans-Enol Equilibrium  
by Means of Infrared Absorption Spectra

SOV/48-22-9-32/40

of acetic ester found by chemical methods. Experimental chemical investigations of a number of keto-enol compounds showed (Ref 1) that the cis-fixed enols accurately adhere to Meyer's law. The equilibrium constant of substances which only exhibit a trans-enol form (trans-fixed enol) is independent of the solvent. Two series of experiments showed that 1) the choice of acetic ester as a standard solvent is justified and 2) that  $L'$  is a constant quantity. Hence it was possible to set up the formula (Ref 7)  $K_T = EL + E_1$ . Keto-enols, in the solution of which cis- and trans-enol variants are contained are also characterized by a linear function of  $K_T$  versus  $L$ . Quantitative measurements of the keto-enol equilibrium which were carried out by chemical methods and by infrared absorption spectra exhibit good accordance. Spectroscopic evidence also validates the general formula for the equilibrium constant of the keto-cis-trans-enol tautomerism. There are 6 figures and 7 references, 3 of which are Soviet.

Card 2/3

Investigation of the Keto-Cis-Trans-Enol Equilibrium  
by Means of Infrared Absorption Spectra

SOV/48-22-9-32/40

ASSOCIATION: Fizicheskiy institut im. P.N.Lebedeva Akademii nauk SSSR  
(Institute of Physics imeni P.N.Lebedev, AS USSR) Komissiya  
po spektroskopii Akademii nauk SSSR (Committee of Spectro-  
scopy, AS USSR) IEOS Akademii nauk SSSR (IEOS, AS USSR)

Card 3/3

SOV/74-27-8-7/7

AUTHOR: Ioffe, S. T. (Moscow)

TITLE: The Vinylation by Means of the Grignard Reaction (Vinilirovaniye posredstvom reaktsii Grin'yara)

PERIODICAL: Uspekhi khimii, 1958, Vol. 27, Nr 8, pp. 1010-1024 (USSR)

ABSTRACT: Among the many types of syntheses using organomagnesium compounds no reactions have hitherto been known permitting the use of the Grignard reagent. Already Krestinskiy showed that under the influence of magnesium on the ether solution of vinyl bromide no organomagnesium compound can be obtained (Refs 1, 2 and 3). The author refers to the Austerweil patent (Austervayl) on the formation of isoprene in the reaction  $\text{CH}_2=\text{CHBr}$  and  $\text{ClC}(\text{CH}_3)=\text{CH}_2$  in the presence of magnesium, and also mentions some other papers (Refs 4 - 14) in order to show that in the practical application of organomagnesium syntheses a gap is found between the Grignard reagent containing saturated halogen alkyls on the one hand and the Iotsich compounds:  $\text{RC}\equiv\text{CMgBr}$  and  $\text{BrMgC}\equiv\text{CMgBr}$  on the other hand (Ref 15). The synthesis of organomagnesium compounds with a vinyl radical under the influence of magnesium upon the halides

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SOV/74-27-8-7/7

## The Vinylation by Means of the Grignard Reaction

$>C = C - X$  (where X is a chlorine or a bromine) in a tetrahydrofuran medium carried out by Normant (Norman) is discussed in detail. The organomagnesium compounds produced containing a vinyl radical (Normant reagents) have the same reactivity as the organomagnesium compounds of halogen alkyls and alkynyls. The production of various unsaturated amino alcohols (Ref 28) proceeding from the amino aldehydes of chlorine hydrines as described by Normant is discussed in detail. In contrast to the reaction of aliphatic Grignard reagents with chloral the Normant reagents with chloral form normal reaction products like the aromatic Grignard reagents: Trichloromethyl-vinyl carbinols. With respect to the elements of the IV. column there are reasons to assume the production of tetravinyl silane under the action of magnesium chloro-vinyl on the tetrachloro silicon in the medium of pentane. The synthesis of the vinyl derivatives of silicon (Ref 36) carried out by Petrov and Mironov is mentioned. The author further mentions that American authors (Ref 39) produced similar substances in good yield also from magnesium chloro-vinyl. Finally he mentions that with respect to the elements of the V. group the description for the production of trivinyl deriva-

Card 2/3

The Vinylation by Means of the Grignard Reaction

SCV/74-27-8-7/7

tives of phosphorus, arsenic, antimony and bismuth from the corresponding halides of the elements may be found in the papers of American authors (Ref 41). The present article is to demonstrate the variety of possibilities of synthesis by employing organomagnesium compounds containing vinyl-groups. A table of the reactions carried out is enclosed. There are 1 table and 41 references, 5 of which are Soviet.

1. Grignard reagents--Synthesis
2. Vinyl compounds--Chemical reactions

Card 3/3

KABACHNIK, M.I.; IOFFE, S.T.; MASTRYUKOVA, T.A.

Tautomerism in aprotic media. Tautomeric equilibrium of phosphorus  
thio acids in benzene and chlorobenzene. Zhur.ob.khim. 30  
no.8:2763-2767 Ag '60. (MIRA 13:8)

1. Institut elementoorganicheskikh soyedineniy Akademii nauk  
SSSR.

(Tautomerism)

(Phosphorus acids)

KABACHNIK, M. I.; IOFFE, S. T.; POPOV, Ye. M.; VATSURO, K. V.

Trans-enolization. Part 1: Effect of solvents on the enolization  
of trans-fixed keto enols. Zhur.ob.khim. 31 no.7:2122-2131 J1 '61.  
(MIRA 14:7)

1. Institut elementoorganicheskikh soedineniy AN SSSR.  
(Ketone) (Isomerism)

KABACHNIK, M.I.; IOFFE, S.T.; POPOV, Ye.M.; VATSURO, K.V.

Transenolization. Part 2: Effect of solvents on the trans-enolization of  $\alpha$ -alkylacetoacetic esters. Zhur.ob.khim. 31 no.8:2682-2692 Ag '61. (MIRA 14:8)

1. Institut elementoorganicheskikh soedineniy AN SSSR.  
(Acetoacetic acid) (Isomerization)

IOFFE, S.T.; POPOV, Ye.M.; VATSURO, K.V.; TULIKOVA, Ye.K.; KARACHNIK, M.I.,  
akademik

Keto cis-trans-enol equilibrium of 3-alkylacetylacetones. Dokl.  
AN SSSR 144 no.4:802-805 Je '62. (MIRA 15:5)

1. Institut elementoorganicheskikh soedineniy AN SSSR.  
(Acetone) (Isomerization)

~~IOFFE~~, Saveliy Timofeyevich; NESMEYANOV, Aleksandr Nikolayevich;  
KOCESHKOV, K.A., otv. red.; OKHLOBYSTIN, O.Yu., red.;  
DOROKHINA, I.N., tekhn. red.

[Methods of the chemistry of organometallic compounds;  
magnesium, beryllium, calcium, strontium, barium] Metody  
elementno-organicheskoi khimii; magnii, berillii, kal'tsii,  
strontsii, barii. Pod obshchei red. A.N.Nesmeianova i K.A.  
Kocheshkova. Moskva, Izd-vo AN SSSR, 1963. 561 p.

(MIRA 16:12)

1. Chlen-korrespondent AN SSSR (for Kocheshkov).  
(Organometallic compounds)

KABACHNIK, M.I.; IOFFE, S.T.

Effect of steric factors on keto-cis-trans-enol equilibrium  
of  $\alpha$ -cycloalkylacetoacetic esters. Izv. AN SSSR. Otd. khim. nauk  
no. 2:339-343 F '63. (MIRA 16:4)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.  
(Acetoacetic acid) (Tautomerism)



L 5064-66 EWT(m)/EPF(c)/ENF(j)/T/ETC(m) RM/DS/WM  
 ACCESSION NR: AP5025507 UR/0062/65/000/009/1556/1564 6/49B  
 541.634+541.67  
 AUTHOR: Molin, Yu. N.; Ioffe, S. T.; Zayev, Ye. Ye.; Solov'yeva, Ye. K.; Kugucheva, Ye. Ye.; Voyevodskiy, V. V.; Kabachnik, M. I.  
 TITLE: Nuclear magnetic resonance study of the keto-enol equilibrium of 3-alkylacetylacetones  
 SOURCE: AN SSSR. Izvestiya. Seriya khimicheskaya, no. 9, 1965, 1556-1564  
 TOPIC TAGS: NMR spectroscopy, ketone, NMR  
 ABSTRACT: NMR spectra of the following compounds were studied: 3-methyl-, 3-ethyl-, 3-n-propyl-, 3-isobutyl-, 3-isopropyl-, and 3-sec-butylacetylacetone, and also 2-isopropoxy-2-penten-4-one. The spectra were taken with a JNM-3 instrument (40 Mc) and some were also recorded with an RS-2 spectrometer (60 Mc) at ~ 25°C, and the content of enol forms was determined. Alkylacetylacetones with unbranched substituents were shown to contain cis-enol forms at equilibrium with the ketone; this agrees with chemical data. Compounds with branched substituents (3-isopropylacetylacetone and 3-sec-butylacetylacetone) were shown to contain only trans-enol forms.  
 Card 1/2

L 5064-66

ACCESSION NR: AP5025507

12  
lacetone) are almost pure ketones. The slight enolization of these substances does not permit the classification of the enol form in the cis or trans series on the basis of the NMR method alone. "Measurements with the RS-2 instrument were made at the Tsentral'nyy institut khimii Vengerskoy Akademii nauk (Central Chemistry Institute of the Hungarian Academy of Sciences) with the direct participation of Dr. L. Radich, to whom the authors express their gratitude." Orig. art. has: 5 figures and 2 tables.

ASSOCIATION: Institut elementoorganicheskikh soedineniy Akademii nauk SSSR (Institute of Organometallic Compounds, Academy of Sciences, SSSR); Institut khimicheskoy kinetiki i goreniya Sibirskogo otdeleniya Akademii nauk SSSR (Institute of Chemical Kinetics and Combustion, Siberian Branch, Academy of Sciences, SSSR)

SUBMITTED: 04Jul63

ENCL: 00

SUB CODE: OC, NP

NO REF SOV: 000

OTHER: 014

Card 2/2 *ML*

KABACHNIK, M.I., akademik; IOFFE, S.T.

Application of correlation equations to keto-enol equilibrium.  
Dokl. AN SSSR 165 no.5:1085-1087 D '65.

(MIRA 19:1)

1. Institut elementoorganicheskikh soedineniy AN SSSR. Submitted July 1, 1965.

IOFFE, S.T.

"The study of tautomerism in aprotic media."

Khimiya i Primeneniye Fosfororganicheskikh Soedineniy (Chemistry and application of organophosphorus compounds) A. YE. APPELOV, Ed.  
Publ. by Kazan Affil. Acad. Sci. USSR, Moscow 1962, 632 pp.

Collection of complete papers presented at the 1965 Kazan Conference on Chemistry of Organophosphorus Compounds.

BALUYEVA, G.A.; IOFFE, S.T.

Beryllium, calcium, strontium, and barium organic compounds.  
Usp.khim. 31 no.8:940-962 Ag '62. (MIRA 15:8)

1. Institut elementoorganicheskikh soyedineniy AN SSSR.  
(Organometallic compounds)

IOFFE, S. Ya.

35591 K voprosu o lechenii vospalitel'nykh zabolevaniy zhenskoy polovoy sfery  
perelivaniyem krovi drcbnymi dozami. Trudy sev-oset. Gos. Med. In-ta, Vyp. 4, 1949,  
C. 69-73 (page 32)

SO: Letopis' Zhurnal'nykh Statey, Vol. 45, 1949

IOFFE, S.Ye., kandidat meditsinskikh nauk

Delivery of twins in uterus bicornis. Akush. i gin. 33 no.2:102-103  
Mr-Apr '57. (MIRA 10:6)

1. Iz kafedry akusherstva i ginekologii (sav. - prof. S.D.Astrinskiy)  
Severo-Osetinskogo meditsinskogo instituta na baze rodit'snogo doma  
No.2 (glavnyy vrach S.G.Dotseva).

(TWINS

in uterus bicornis, normal delivery)

(DELIVERY

twins, in uterus bicornis)

SHEMET, Aleksey Savel'yevich; IOFFE, S.Ye., redaktor; SAVICH, M.P., redaktor;  
OYSTRAKH, V.G., tekhnicheskiy redaktor

[Work practice of a crew of conveyor movers] Opyt raboty kompleksnoi  
brigady posadoperenoschikov, Alma-Ata, Kazakhskoe gos. izd-vo,  
1956. 13 p. (MLRA 9:10)

1. Brigadir kompleksnoy brigady posadoperenoschikov uchastka No. 3  
shakhty No.35 kombinata "Karagandaugol" (for Shemet)  
(Mine haulage)



TAKIROV, Musatay; IOFFE, S. Ye., redaktor; ZAV'YALOV, G.P., redaktor;  
OYSTRAKH, V.G., tekhnicheskii redaktor

[The party group in the struggle for technical progress] Partinaiia  
gruppa v bor'be za tekhnicheskii progress. Alma-Ata, Kazakhskoe gos.  
izd-vo, 1956. 18 p. (MLRA 9:10)

1, Mashinist ugol'nogo kombayna, partgruppovaya shakhty No.1-bis tresta  
Kirovugol' kombinata "Karagandaugol'" (for Makirov)  
(Communist Party of the Soviet Union--Party work)  
(Coal mines and mining)

KULAKOV, Nikolay Pavlovich; ~~LOVE~~ S. Ya., redaktor; ZAV'YALOV, G.P.,  
redaktor; OYSTRAKH, V.G., tekhnicheskiy redaktor

[The communists are the organizers of the struggle for coal; the  
work practice of the Party group in the Kalinin Mine of the  
"Karagandaugol'" combine] Kommunisty - organizatory bor'by za ugol';  
kombinata "Karagandaugol'". Alma-Ata, Kazakhskoe gos. izd-vo, 1956.  
14 p. (MIRA 9:10)

(Communist Party of the Soviet Union--Party work)  
(Coal mines and mining)

IOFFE, Solomon Yefimovich; VAL'SHTEYN, G., redaktor; DOLGOPIYATOV, Yu.,  
redaktor; OYSTRAKH, V., tekhnicheskii redaktor

[General combined work crews in mines] Skvoznye kompleksnye brigady  
na shakhte. Alma-Ata, Kazakhskoe gos. izd-vo, 1956. 13 p. (MLRA 9:10)  
(Coal mines and mining)

IOFFE, T.P.

Treatment of hypertension with an infusion of eucalyptus under  
polyclinical conditions. Sov. med. 24 no.6:106-107 Je '60.  
(MIRA 13:9)

1. Iz polikliniki No 7 Kuybyshevskogo rayona Moskvyy.  
(HYPERTENSION) (EUCALYPTUS)

IOFFE, TS. I.

32412, Ioffe, Ts. I. K metodike izucheniya snosa benticheskiykh organizmov rekoy i yego rol' v zaselenii vodokranilishcha. Izvestiya Vsesoyuz. nauch.-issled. in-ta ozer. irech. ryb. khoz-va, t. XXIX, 1949, s. 75-95. --- Bibliogr: 12 nazv.

SO: Letopis' Zhurnal'nykh Statey, Vol. 44

IOFFE, TS.I.

Formation of benthonic fauna of Rybinsk Reservoir. Trudy probl.  
1 tem.soveshch. no.2:32-40 '54. (MIRA 8:5)  
(Rybinsk Reservoir--Fresh-water fauna)

IOFFE, T.S.I.

Increase of food reserves in ponds by means of organic fertilizers.  
Trudy probl. i tem. soveshch. no.2:112-121 '54. (MLRA 8:5)  
(Fish ponds)

Fogge, Ts. I.

AG <sup>10</sup> Increasing the nutrition base of fish ponds with organic  
fertilizer. Ts. I. Ioffe. *Trudy Problemy i Temy. Seriya  
Ishkaniy, No. 2, Priklady Gidrobiol. Vnutren. Vod No. 2,  
112-21(1954).*—Horse manure, branches of alder, mis-  
cellaneous aquatic plants, and miscellaneous weeds were  
added to different ponds. The alder branches caused a  
higher reduction of  $O_2$  than any other org. material, reduced  
the water, and increased the acidity, quantity of  $NH_4$  salts,  
and of sol. Fe. Weeds were next in their effect, and manure  
was least of all treatments. J. S. Joffe



IOFFE, TS.I.

Effect of fertilizers on the increase in the food supply of ponds.  
Trudy probl. i tem. sov. no.7:68-72 '57. (MLRA 10:4)  
(Fish ponds) (Fertilizers and manures) (Fishes--Food)

IOFFE, TS.M., starshiy nauchnyy sotrudnik

Behcet disease; personal observations. Vest. oft. 71 no.2:16-22  
Mr-Ap '58. (MIRA 11:4)

1. Institut glaznykh bolezney imeni Gel'mgol'tsa (dir.-kand.med.nauk  
A.V. Roslavytsev)  
(BEHCET SYNDROME, case reports)

IOFFE, V.

PA 51/49T92

USSR/Radio  
Radio Receivers

Jan 49

"Requirements for a Radio Receiver," V. Ioffe,  
A. Godzevskiy, 3 pp

"Radio" No 6

Discusses A. Frolov's article which listed re-  
quirements for a high-quality receiver. Authors  
take exception to Frolov's conclusions on fre-  
quency response range for the speaker, maximum  
nonlinear distortion permissible, output power,  
number of bands, etc.

51/49T92

*Ioffe, V.*

4-11-33/34

AUTHOR: Ioffe, V., Engineer

TITLE: A Metallurgist's Notes (Zametki metallurga)

PERIODICAL: Znaniye - Sila, 1957, # 11, p 47 (USSR)

ABSTRACT: Under the heading "The Earth's Treasures" the article states that the Earth's crust contains approximately 775,000,000 billion tons of pure iron. The quantity of aluminum in the Earth's crust exceeds 1,370,000,000 billion tons and that of copper 2,000,000 billion tons, gold - 93.5 billion tons.

The next part of the article is entitled "The World's Output of Metals" and contains particulars about the production of cast iron, copper, aluminum, etc. at different times. The last part deals with the cost of 1 ton of metal, and contains comparative figures in regard to the quantity of electric power used for the production of various metals.

There are 6 figures.

AVAILABLE: Library of Congress

Card 1/1

*IOFFE V.*  
ZENKOVICH, V., prof. doktor geogr. nauk; LAGUNOVA, I.; PETROVSKIY, Yu.  
zhurnalist; VERD'YE, Zhan; PETROV, S., inzh.; NAUMOV, S., nauchnyy  
soтрудnik; IOFFE, V., inzh.; DROZDOV, V., inzh.

People of new specialties. Znan. sila 32 no.11:32-34 W '57.

(MLBA 10:11)

1. Direktor Instituta rentgenologii i radiologii Ministerstva zdrazvo-  
okhraneniya (for Lagunova)

(Science)

IOFFE, V.

Greatest document of our time. Plast.massy no.10:1-2 '61.  
(Plastics) (MIRA 15:1)

IOFFE, V.

Paradoxes in insulation. Znan.-sila 37 no.6:46 Je '62.  
(Electric insulators and insulation) (MIRA 15:9)

10

**10FFE, V.A.**

On the acid. V. A. Ioffe and A. V. Vostainskaya.  
 Russ. 31,430, Oct. 31, 1953.  $\text{Na}_2\text{C}_2\text{O}_4$  is treated with  
 $\text{H}_2\text{SO}_4$  to ppt.  $\text{NaHC}_2\text{O}_4$ . This is filtered off and further  
 treated with  $\text{H}_2\text{SO}_4$  to form  $\text{H}_2\text{C}_2\text{O}_4$ .

ASB-5.4 METALLURGICAL LITERATURE CLASSIFICATION

FROM STUDENT

EXAMINED BY

DATE

REMARKS

10FFE, V.A.



1ST AND 2ND SECTIONS										PROCESSES AND PROPERTIES INDEX									
<p><b>IOFFE V. A.</b></p> <p><i>CC</i></p>										<p>11. 21.</p> <p>The Kerr effect in solution. V. A. Ioffe. <i>Compt. rend. acad. sci. (U. R. S. S.)</i> 3, 401-3 (in German, 1934-35) (1934).—The Kerr consists of mins. of nitrobenzene in Ccl<sub>4</sub>, CCl<sub>2</sub> and heptane were detd. At low concn. of nitrobenzene the const. does not change, but at a definite concn. it begins to decrease rapidly, approaching the value for pure liquid nitrobenzene. The behavior is connected with assocn. of nitrobenzene. A. B. P. Duncan</p>									
<p>ASB-55.8 METALLURGICAL LITERATURE CLASSIFICATION</p>										<p>HOW BORING</p>									
<p>10000 21000 30000 40000 50000 60000 70000 80000 90000</p>										<p>10000 20000 30000 40000 50000 60000 70000 80000 90000</p>									

**OFFE, V A**

**BC**

Raman spectra of binary silicate glasses.  
M. F. Vukobratovic and V. A. Jovanovic (Bull. Acad. Sci. U.R.S.S.,  
1938, Cl. Sci. Tech., No. 7, 672-680).—Na glasses have  
4, and Pb glasses 2, characteristic Raman lines, not  
content the spectra gradually approach that of SiO<sub>2</sub>.  
Na glass has a very intense line 940–960 cm<sup>-1</sup>,  
which gradually fades as the SiO<sub>2</sub> content rises, and in  
ascribed to Na<sub>2</sub>SiO<sub>3</sub>. The shift in the spectrum ob-  
served as the Pb content is progressively raised corre-  
sponds with substitution of Na by Pb atoms. R. T.

Dr. abs. IOFFE, V. A.

Bl-1, Chemical Engineering, Plant,  
Machinery. (Unit Processes).

Evaporation of liquids in closed vessels. M. V. A. Ioffe, M. V. Kovalchuk, and M. M. Babayev. *J. Tech. Phys. USSR*, 1968, 38, 3, 341-344. An equation was derived for the evaporation of a liquid from the bottom of a vertical cylindrical tube originally filled with pure gas. The process is treated in the first stage according to the equation:  $\frac{dP}{dt} = \frac{P_0 - P}{\tau} \ln \frac{P_0}{P}$ , where  $P$  is the pressure in atm. at beginning of evaporation,  $P_0$  is the initial atm. in atm. per sec.,  $t$  is the time in sec.,  $\tau$  is the evaporation time in sec. and  $V$  is the vol. of tube in cc. This equation is used for evaporation of benzene in air in a closed vessel, and for  $CCl_4$  in SF<sub>6</sub>. The apparatus is described and in both cases the evaporation is found to follow the above equation closely in the initial stages. W. H. H. H.

Lab. Silicate Chem., AS USSR

10PPE, U. H.

41 Temperature dependence of the dielectric loss in two-component silicate glass systems ( $K_2O-SiO_2$  and  $PbO-SiO_2$ ). V. A. Iskh. Doklady Akad. Nauk S.S.S.R. 27, 405-8 (1942). The dielec. loss of  $K_2O-SiO_2$  systems (contg. 17.3-22.7 mol. %  $K_2O$ ) and of  $PbO-SiO_2$  systems (14.3-59.3 mol. %  $PbO$ ) was studied for the temp. range 12 to 420°K. at a frequency of 1 Mc. For the potassium silicate systems, the max. of dielec. loss occur below 100°K. and the min. occur between 130 and 160°K. Above 300°K., the dielec. loss increases with temp. in an exponential manner. For lead silicate glass systems as well as lead silicate glass contg.  $TiO_2$ , there is a max. of dielec. loss at about 100°K. but no increase of the loss with temp. for the range studied. The following explanations are given. At low temp., there is a decrease of the dielec. loss with temp. since the loss is due to deformation of the silicon-oxygen net work. As the temp. increases, the mobility of the  $K^+$  ions becomes significant and the dielec. loss in potassium silicate systems increases. For lead silicates, the mobility of  $Pb$  ions remained unimportant for all temps. studied, and consequently no increase in dielec. loss is observed with increasing temp. Paul V. Fing.

IOFFE, V. A.

FD-434

USSR/Physics - Dielectric losses

Card 1/1 : Pub. 153 - 4/18

Author : Ioffe, V. A.

Title : Dielectric losses in silicate glasses

Periodical : Zhur. tekhn. fiz. 24, 611-621, Apr 1954

Abstract : Investigates the temperature dependence of the dielectric losses of glasses of the systems  $K_2O-SiO_2$ ,  $PbO-SiO_2$ ,  $Na_2O-SiO_2$ , etc. in the temperature range 120 to 4200 K at a frequency of  $10^6$  cycles. Finds that for low temperatures the dielectric losses of all glasses studied possess a maximum that shifts at higher frequencies towards higher temperatures. Thanks Prof. A. I. Shal'nikov and G. A. Smolenskiy. Briefly surveys the literature. 28 references, including 12 Soviet.

Institution : —

Submitted : June 23, 1953

IOFFE, V. A.

USSR/Chemical Technology. Chemical Products and Their Application -- Silicates.  
Glass. Ceramics. Binders, I-9

Abst Journal: Referat Zhur - Khimiya, No 2, 1957, 5182

Author: Ioffe, V. A.

Institution: Academy of Sciences USSR

Title: Dielectric Losses in Silicate Glasses

Original

Publication: Sb. Stroyeniye stekla, M.-L., AN SSSR, 1955, 258-263

Abstract: Description of the results of investigations of dielectric losses of two-component potassium-, lead- and sodium silicate glasses, and also of glasses containing magnesium ions. It was found that losses in lead glasses are almost not dependent on the composition; the same is also observed in potassium glasses, and only in the sodium glasses the losses increase appreciably with increase in  $\text{Na}_2\text{O}$  content. In the region of medium and low frequencies the dependence of losses on composition is very great, especially in the case of alkali glasses; losses increase rapidly with increasing content of alkali ions.

Card 1/1

Ioffe, V.A.

USSR/Chemical Technology. Chemical Products and their Application.  
Glass. Ceramics. Building Materials.

J-12

Abs Jour: Referat Zh.-Kh., No 8, 1957, 27627

Author : V.A. Ioffe.

Inst :

Title : Information.

Orig Pub: vSb: Stroyeniye stekla. M.-L., AN SSSR, 1955, 327.

Abstract: The author repudiates his former assertions that sodium bisilicate is found in sodium-silica glass. See also RZhKhim., 1957, 5166 and 5182.

Card : 1/1

-28-

IOFFE, V. A.

USSR/Chemical Technology. Chemical Products and Their Application -- Silicates.  
Glass. Ceramics. Binders, I-9

Abst Journal: Referat Zhur - Khimiya, No 2, 1957, 5183

Author: Ioffe, V. A.

Institution: Academy of Sciences USSR

Title: Reply to G. I. Skanavi

Original

Publication: Sb. Stroyeniye stekla, M.-L., AN SSSR, 1955, 342-343

Abstract: It is pointed out that the neutralization effect manifests itself, in alkali glasses, not only as concerns dielectric losses and electric conductivity, but also by a lowering of the coefficient of linear expansion, increase in chemical stability, etc. This is not only indicative of the fact that alkali ions are bound stronger, but also that the silicon-oxygen network is more rigid in such glasses.

Card 1/1



IOFFE, V. A.

G-2

Category : USSR/Electricity - Dielectrics

Abs Jour : Ref Zhur - Fizika, No 1, 1957 No 1516

Author : Ioffe, V.A.

Title : Dielectric Losses in Alkali-Borate Glass at Low Temperatures

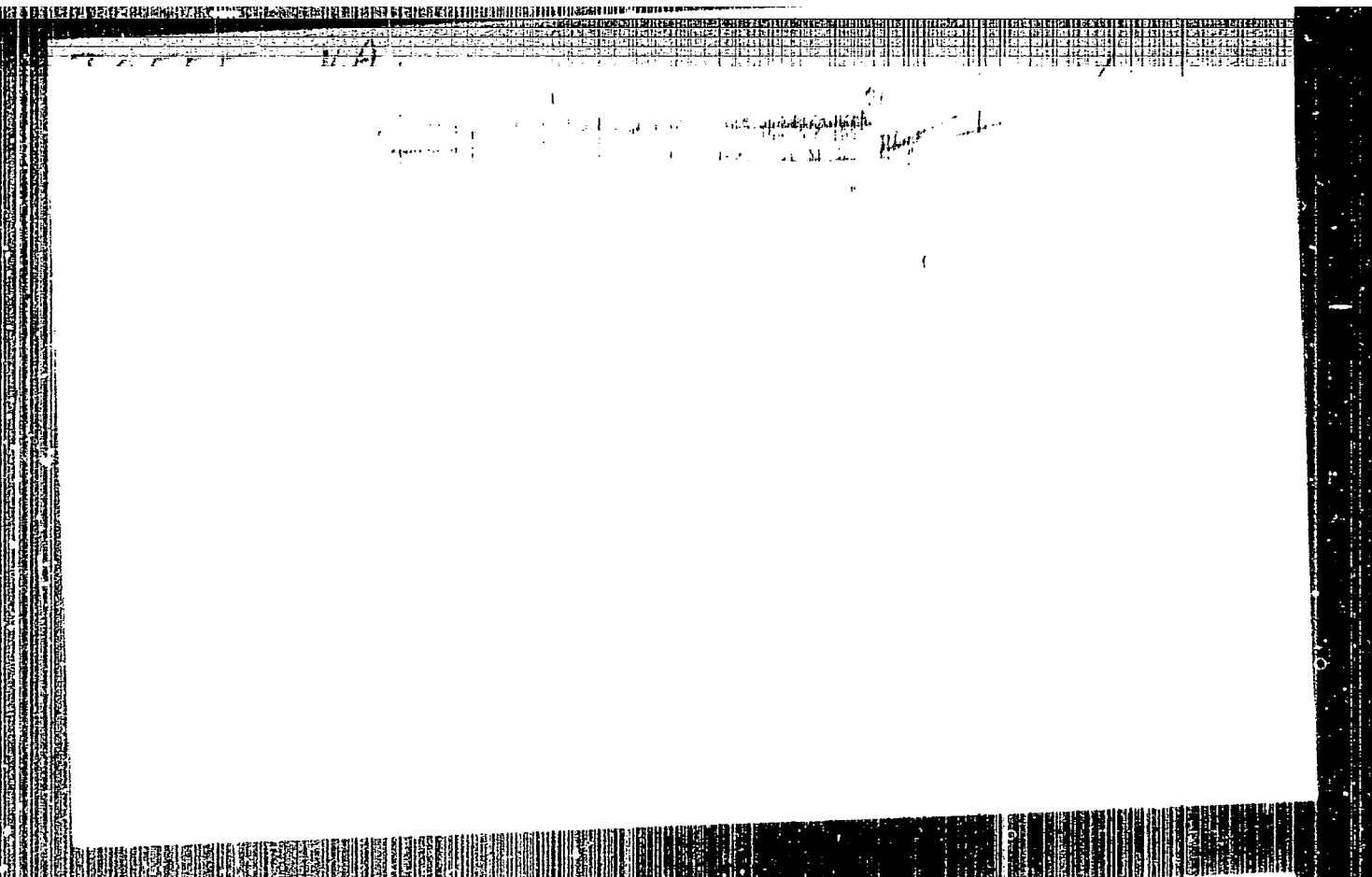
Orig Pub : Zh. tekhn. fiziki, 1956, 26, No 3, 516-525

Abstract : The dielectric loss angle of alkali-borate glasses of various compositions was measured in the 12 -- 3000 K range at frequencies of  $2.4 \times 10^2$  --  $10^6$  cycles. The  $\tan \delta$  vs. T curve exhibits a maximum, the magnitude and position of which depends on the contents of the alkali oxide in the glass. In the author's opinion, the dielectric losses at low temperatures are caused by the relaxation of the elements of the structural grid of the glass. At room temperatures the activation energy of the process becomes comparable with the thermal-motion energy and the losses are caused by the anharmonic-oscillatory motion of the structural elements. Upon mutual replacement of the alkali cations, the neutralization effect is observed when their total concentration reaches 25 molar percent and above.

Card : 1/1

"APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000618630003-4



APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000618630003-4"

Ioffe, V. A.

57-27-7-7/40

AUTHOR: Ioffe, V. A.

TITLE: On the Nature of the Dielectric Losses in Sodium-Aluminum-Silicate Glasses (O prirode dielektricheskikh poter' v natrevoaluminosilikatnykh steklakh)

PERIODICAL: Zhurnal Tekhnicheskoy Fiziki, 1957, Vol. 27, Nr 7, pp. 1454 - 1461 (USSR)

ABSTRACT: The dependence of the dielectric losses on the oxygen number (polymerization-degree of the structural network) in the range of from 20 to 290° K at frequencies of  $10^6$ ,  $2,3 \cdot 10^5$  and  $1,7 \cdot 10^6$  cycles is investigated. It is shown that the dielectric losses increase with a decrease in the oxygen number in all glasses investigated (feldspar  $1 \text{ Na}_2\text{O} \cdot 1 \text{ Al}_2\text{O}_3 \cdot 6 \text{ SiO}_2$ , jade  $1 \text{ Na}_2\text{O} \cdot 1 \text{ Al}_2\text{O}_3 \cdot 4 \text{ SiO}_2$ , nephelite  $1 \text{ Na}_2\text{O} \cdot 1 \text{ Al}_2\text{O}_3 \cdot 2 \text{ SiO}_2$ , an intermediate product  $1 \text{ Na}_2\text{O} \cdot 1 \text{ Al}_2\text{O}_3 \cdot 3 \text{ SiO}_2$ , besides two glasses with an oxygen number below two were welded together:  $1 \text{ Na}_2\text{O} \cdot 1 \text{ Al}_2\text{O}_3 \cdot 3,9 \text{ SiO}_2$  and  $1 \text{ Na}_2\text{O} \cdot 1,2 \text{ Al}_2\text{O}_3 \cdot 3,8 \text{ SiO}_2$ ). It is shown that at room temperature the angle of the dielectric losses in glasses with equal oxygen number increases with increasing sodium-oxide content in the glass. In the range of low temperatures the angle is not dependent on the sodium-oxide content, but increases with increasing aluminum-oxide content

Card 1/3

57-27-7-7/40

On the Nature of the Dielectric Losses in Sodium-Aluminum-Silicate Glasses

and with a decrease in the oxygen number. The supposition on the reconstruction of the glass-network and the transition of aluminum into the position of a modifier could not be considered well-founded in any of the glasses investigated. In the sodium-aluminum-silicates aluminum is in a tetrahedron-coordination, independent of the structure of the corresponding crystalline sodium-aluminum-silicate. The conception at present existing on the structure of the dielectric losses in the glasses are not sufficient for an explanation of the high value of the phase angle (loss angle) in sodium-aluminum-silicate glasses and its dependence on the oxygen number. The supposition is expressed that the displacements connected with the negative aluminum-oxygen-tetrahedrons of the holes form an additional source of loss in these glasses. It is to be expected that the electric properties beside the ion- and electron-processes can be determined in glasses which in the structural network possess ions with a valence different from four that isomorphously replace the silicon-ions. There are 8 figures, 1 table and 32 references, 11 of which are Soviet.

Card 2/3

57-27-7-7/40  
On the Nature of the Dielectric Losses in Sodium-Aluminum-Silicate Glasses

ASSOCIATION: Institute for the Chemistry of Silicates AS USSR, Leningrad  
(Institut khimii silikatov AN SSSR, Leningrad)

SUBMITTED: February 6, 1957

AVAILABLE: Library of Congress

1. Glass-Dielectric properties

Card 3/3

IOFFE, V.A.

57 - 9-10/40

AUTHORS

Ioffe, V.A., Khvostenko, G.I.,  
Zonn, Z.N.

TITLE

The Electrical Properties of Some Single Crystals and  
Polycrystalline Ferrites.  
(Elektricheskiye svoystva nekotorykh monokristallov i  
polikristallicheskikh ferritov.)

PERIODICAL

Zhurnal Tekhn. Fiz., 1957, Vol. 27, Nr 9, pp.1985-1995  
(USSR)

ABSTRACT

The dependence of the specific resistance, of the  
dielectricity constant, and of the angle of dielectric  
losses on temperature at sound frequencies and for solid  
solutions of nickel-ferrite and zinc-ferrite, of  
magnesium-ferrite and manganese-ferrite as well as in the  
case of two single crystals and a ceramic sample of a  
solid solution of cobalt-ferrite and zinc-ferrite was  
investigated. All ferrites investigated have a high  
dielectricity constant within the range of low frequencies  
and high temperatures. The dependence of the dielectricity  
constant on frequency and temperature is due to relaxation  
processes. It is shown that the dielectricity constant of  
ferrites is a property that is independent of their poly-

CARD 1/2

57-9-10/40

The Electrical Properties of Some Single Crystals and Polycrystalline Ferrites.

crystalline structure. The presence of ions of one and the same with different valence in the ferrites, in which case the ferrites are distributed statistically in equal crystallographic positions, cause - apart from through-conductivity - the occurrence of electron displacements under the influence of the field. These local displacements cause high electron polarisation in the ferrites.

There are 15 figures, 2 tables and 2 Slavic references.

ASSOCIATION:

Institute for the Chemistry of Silicates AN USSR,  
Leningrad.

SUBMITTED:

(Institut khimii silikatov AN SSSR, Leningrad.)

AVAILABLE:

March 25, 1957  
Library of Congress.

CARD 2/2

SOV/1-2-10-9/40

24(6)

AUTHORS:

Ioffe, V. A., Yanchevskaya, I. S.

TITLE:

Dielectric Losses in Feldspars (Dielektricheskiye poteri v polevykh shpatakh)

PERIODICAL:

Zhurnal tekhnicheskoy fiziki, Vol 28, Nr 10, pp 2154-2164 (USSR), 1978

ABSTRACT:

This is an investigation of the temperature and frequency dependence of the loss angle  $\tan \delta$  and of the dielectric constant  $\epsilon$  of a number of natural monocystals of feldspars in the temperature range of 20 - 500°K and a frequency region of  $5 \cdot 10^2$  -  $5 \cdot 10^6$  cy. This paper covers the isomorphic series of sodium-potassium feldspars, the plagioclases (which are a continuous series of solid solutions of albite ( $\text{NaAlSi}_3\text{O}_8$ ) with anorthite ( $\text{CaAl}_2\text{Si}_2\text{O}_8$ ), and microcline ( $\text{KAlSi}_3\text{O}_8$ ). A resonance absorption and an anomalous dispersion of the dielectric constant was found to exist in all feldspars investigated at a frequency of  $5 \cdot 10^5$  cps. In the range of 200 - 500°K the dielectric losses in feldspars are caused by resonance phenomena, resonance occurring

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Dielectric Losses in Feldspars

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by thermal excitation. If the temperature is raised, the resonance frequency also rises at first. This explains the existence of the maxima in the  $\text{tg } \delta$  versus temperature function at sonic frequencies. In microline the  $\text{tg } \delta$  and  $\epsilon$  versus frequency functions exhibit two maxima and two domains with an anomalous dispersion of  $\epsilon$ . As the resonance frequency observed in all feldspars is low and identical this resonance must necessarily be ascribed to electron processes. There is every indication that the resonance is due to the transition of an electron from one oxygen atom into another oxygen atom in the negatively charged aluminum-oxygen tetrahedron. This conception is, however, of a still preliminary nature. There are 21 figures, 1 table, and 8 references, 4 of which are Soviet.

SUBMITTED: March 21, 1958

Card 2/2

AUTHORS: Ioffe, V. A., Khvostenko, G. I.

20-118-4-23/61

TITLE: The Anomalous Dispersion of the Dielectric Constant in Feldspars (Anomal'naya dispersiya dielektricheskoy pronitsayemosti v polevykh shpatakh)

PERIODICAL: Doklady Akademii Nauk SSSR, 1958, Vol. 118, Nr 4, pp. 709-712 (USSR)

ABSTRACT: At first, the authors shortly report on the development of this problem. The present paper investigates the dielectric loss angle and the dielectric constant of potassium feldspar (orthoklase) and of sodium feldspar (albite) within the temperature range of from 20 to 500°K and within the range of frequencies of from 500 kilocycles to 5 Megacycles. The measurements were performed in vacuo, after the sample was heated to 500°K in an evacuated bell. The electrodes were applied by means of burning-in a silver paste. Measurements were conducted with a bridge circuit. A diagram illustrates the temperature dependence of  $\tan \delta$  and  $\epsilon$  in orthoklase for frequencies of  $8 \cdot 10^5$  and  $8 \cdot 10^4$  cycles. The value of  $\tan \delta$  is very small in orthoklase

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The Anomalous Dispersion of the Dielectric Constant in Feldspars 20-118-4-23/61

at temperatures of from 20 to 500°K ( $\sim 5 \cdot 10^{-4}$ ) and is little dependent upon temperature. - The dielectric constant retains its constant value of  $\epsilon = 6$ . From 200°K onwards  $\text{tg } \delta$  and  $\epsilon$  begin to increase sharply with growing temperature,  $\text{tg } \delta$  increasing by about two orders of magnitude. If the temperature is further raised,  $\text{tg } \delta$  remains constant. A sharp increase of  $\epsilon$  is also observed within the same temperature range. A second diagram illustrates the frequency dependence of  $\text{tg } \delta$  and  $\epsilon$  in orthoklase at the temperatures 297°K, 399°K and 246°K. The maximum of  $\text{tg } \delta$  at all three temperatures is found at the frequency  $\sim 4,5 \cdot 10^5$  cycles. The frequency of the maximum is independent from temperature. A second, wider maximum is observed at a frequency of  $2 \cdot 10^6$  cycles at a temperature of 297°K. Further numerical data are given. The dielectric constant decreases within the range of low frequencies at all temperatures investigated, when the frequency is increased, then passes through a maximum at the frequency of  $2 \cdot 10^5$  cycles, and through a low minimum at  $4,5 \cdot 10^5$  cycles. Then the dielectric constant increases again

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The Anomalous Dispersion of the Dielectric Constant in Feldspars 20-118-4-23/61

up to a value of  $\sim 6,3 \cdot 10^5$  cycles. A further diagram illustrates the temperature dependence of  $\text{tg } \delta$  and of  $\epsilon$  in albite at the frequencies  $8 \cdot 10^5$  and  $8 \cdot 10^4$  cycles. This temperature dependence shows the same character as in orthoklase. Similar dependences were also obtained by the authors for plagioclase, which consists of a solid solution of sodium- and potassium feldspars. An anomalous dispersion of  $\epsilon$  also exists in plagioclase, the range of dispersion, however, is somewhat lower, within the frequency range of  $\sim 10^5$  cycles. The here obtained temperature dependences of  $\text{tg } \delta$  and of  $\epsilon$  in feldspars can neither be explained by conduction processes, nor by relaxation processes. This also holds for the temperature dependence of  $\text{tg } \delta$ . Such a temperature dependence can obviously be explained by resonance phenomena. The resonance phenomena observed in feldspars are obviously caused by electron processes. There are 4 figures, and 2 references, 1 of which is Soviet.

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The Anomalous Dispersion of the Dielectric Constant in Feldspars 20-118-4-23/61

ASSOCIATION: Institut khimii silikatov Akademii nauk SSSR  
(Institute for Silicate Chemistry, AS USSR)

PRESENTED: August 19, 1957, by A. F. Ioffe, Member of the Academy

SUBMITTED: August 16, 1957

AVAILABLE: Library of Congress

Card 4/4

IOFFE, V.A. [translator]; SMOLENSKIY, G.A., red.; FUKTSEV, A.K., red.;  
KORNILOV, B.I., tekhn.red.; POTAPENKOVA, Ye.B., tekhn.red.

[Dielectric spectroscopy; recent studies on the properties of  
certain ferromagnetic semiconductors and dielectrics: relaxation  
processes, electric conductance, losses, and the role of structural  
defects. Translated articles] Dielektricheskaya spektroskopiya;  
noveishie issledovaniya svoystv nekotorykh ferromagnitnykh polu-  
provodnikov i dielektrikov: relaksatsionnyye protsessy, elektropro-  
vodnost', poteri i rol' defektov struktury. Sbornik statei. Pod  
red. G.A.Smolenskogo. Moskva, Izd-vo inostr.lit-ry, 1960. 362 p.  
(MIRA 14:4)

(Spectrum analysis) (Dielectrics) (Semiconductors)



81364

S/181/60/002/03/22/028  
B006/B01715.2120  
24.7700

AUTHORS: Ioffe, V. A., Khvostenko, G. I.

TITLE: Electrical Conductivity of Sodium-aluminum-silicate Glasses

PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 3, pp. 509-516

TEXT: The authors investigated the electrical conductivity<sup>21</sup> of glasses of the system  $\text{Na}_2\text{O} \cdot x \text{Al}_2\text{O}_3 \cdot (y-x) \text{SiO}_2$  with  $y = 2, 3, 4$ , and 6 and  $x$  from 0 to 1.1. The aim of the present investigations was to find out whether a second type of charge carrier exists in these glasses (it has been assumed already earlier that the electrical properties of these glasses are not only determined by ionic but also by electronic processes). They also wanted to investigate the dependence of conductivity on the  $\text{Na}_2\text{O}$  content and the structure at very low temperatures and in the range 15 - 240°C. The composition of the glasses investigated is given in a Table (p. 510). Conductivity was measured electrometrically (Fig. 1), the apparatus made it possible to measure currents of down to  $10^{-14}\text{A}$ ; voltage sensitivity was  $10^{-3}\text{V/graduation}$ . All measurements were made in vacuum,

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Electrical Conductivity of Sodium-aluminum-silicate Glasses

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B006/B017

after a continuous heating of the sample at 250°. Since the initial amperage could not be measured, the time dependences of "charge" and "discharge" of the sample were determined and then extrapolated for  $t = 0$  both graphically as well as by computation. Figs. 2 - 5 show  $\log \sigma = f(1/T)$  of four series of glasses. The following results were obtained: The electrical conductivity in the glasses investigated does not depend on the  $\text{Na}_2\text{O}$  content; it is determined by the ratio between the number of the aluminum-oxygen tetrahedra and the number of silicon-oxygen tetrahedra in the structural lattice, i.e., by  $\text{Al/Si}$ . With increasing  $\text{Al/Si}$ , electrical conductivity increases, whereas the activation energy  $U$  and the number of carriers decreases. The electrical conductivity of two glasses may be expressed by the formula  $\sigma = \sigma_0 \exp(-U_1/kT) + \sigma'_0 \exp(-U_2/kT)$  which indicates that in these glasses two types of carrier exist. The authors assume that in the second type electrons are concerned. The resulting dependence of  $\sigma$ ,  $U$ , and  $\sigma_0$  on the composition (Figs. 7 - 10) may be explained by a change of the ratio of the fractions of ionic and electronic conduction in these glasses. N. M. Verebeychik and V. I. Odilevskiy are mentioned. There are 10 figures, 1 table, and 4 references:

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Electrical Conductivity of Sodium-aluminum-  
silicate Glasses

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B006/B017

3 Soviet and 1 Swiss.

ASSOCIATION: Institut khimii silikatov AN SSSR Leningrad (Institute of  
Silicate Chemistry of the AS USSR, Leningrad)

SUBMITTED: June 4, 1959

Card 3/3

IOFFE, V. A.

81957  
S/181/60/002/04/17/034  
B002/B06324,2100  
AUTHORS:Ioffe, V. A., Patrina, I. B., Poberovskaya, I. S.

TITLE:

Electric Properties of Some Semiconducting Oxide Glasses

PERIODICAL: Fizika tverdogo tela, 1960, Vol. 2, No. 4, pp. 656-662

TEXT: The authors examined glasses of the systems  $V_2O_5 - P_2O_5$ ,  $V_2O_5 - P_2O_5 - BaO$ , and  $WO_3 - P_2O_5 - K_2O$  (Table). Their electrical conductivity  $\sigma$  was measured by means of a tube voltmeter having a  $1 \div 10^{11} \Omega$  tube. A Q-meter of the type Тесла В211 (Tesla V211) and a bridge of the type Тесла М-351 (Tesla M-351) were used to measure the dielectric losses ( $\tan \delta$ ) and the dielectric constant  $\epsilon$ .  $\sigma$  was determined between 290 K and 500 K. Figs. 1, 2, and 6 show the temperature dependence of  $\sigma$  for the above-mentioned systems. The electrical conductivity of the glasses rises with their content of vanadium and tungsten. This is due to the fact that the conductivity is effected by electron transition between vanadium and tungsten ions of different valences. The conductivity of vanadium glasses mainly depends on the ratio of vanadium oxide to phosphorus oxide, and is

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Electric Properties of Some Semiconducting  
Oxide Glasses

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B002/B063

independent of their content of barium (Fig. 2) or sodium (Fig. 9 ; the values were derived from Ref. 5).  $\tan \delta$  and  $\epsilon$  were measured at 1, 50, 500 kc/s, and 1.6 Mc/s within the temperature range 60 - 400°K (Figs. 4, 5, and 8). Here, relaxation processes occur, which are connected with electron transitions between ions of different valences, and are caused by the statistically disordered distribution of these ions. Mention is made of I. I. Kitaygorodskiy and V. G. Karpechenko. There are 9 figures, 1 table, and 8 references: 2 Soviet, 3 American, 2 British, and 1 Czech.

ASSOCIATION: Institut khimii silikatov AN SSSR, Leningrad  
(Institute of Silicate Chemistry of the AS USSR, Leningrad)

SUBMITTED: August 1, 1959

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24934

S/181/61/003/006/031/031  
B102/B21415.2120

AUTHORS: Ioffe, V. A. and Zonn, Z. N.

TITLE: Glasses with high dielectric constant

PERIODICAL: Fizika tverdogo tela, v. 3, no. 6, 1961, 1902-1904

TEXT: The authors investigated the possibility of making glasses on a  $\text{Bi}_2\text{O}_3$ -basis by adding  $\text{TiO}_2$ ,  $\text{BaO}$ , or  $\text{PbO}$  without using a vitrifying oxide (such as  $\text{SiO}_2$ ,  $\text{B}_2\text{O}_3$ , or  $\text{P}_2\text{O}_5$ ). With 70-80 mole%  $\text{Bi}_2\text{O}_3$  and addition of 20-30 mole% at 1100-1150°C a transparent melt was obtained which crystallized when cast into molds. The material had an  $\epsilon$  of 75-80. In the range 100 cps - 2.5 Mc/sec  $\epsilon$  was independent of frequency.  $\tan\delta = 0.005 - 0.006$  at 100 cps, and 0.01 at 2.5 Mc/sec. On addition of 10 mole%  $\text{SiO}_2$  or  $\text{B}_2\text{O}_3$  transparent glasses were obtained which, however, still had the tendency to crystallize and devitrification. Stable glasses were obtained only after adding  $\text{SiO}_2$  in quantities of over 15 mole%. The composition of the glasses investigated are shown in Fig. 1.

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S/181/61/003/006/031/031  
B102/B214

## Glasses with high dielectric constant

X

The glasses were founded in corundum and platinum crucibles in a silite furnace for 30 min. Longer holding times led to a darkening of the glass and a tendency towards devitrification. The softening point of the glasses was 550-850°C. They were stable against water. The glasses with 17-25 mole%  $\text{SiO}_2$  had an  $\epsilon$  of 38-40 which decreased with increasing  $\text{SiO}_2$  content (40 mole%  $\text{SiO}_2$ ,  $\epsilon = 25$ ).  $\epsilon$  was independent of frequency in the range 100 cps-2.5 Mc/sec, and increased linearly with increasing temperature;  $\tan\delta = 0.002-0.003$  at 100 cps, and was only slightly frequency dependent. The crystallization of the glasses led to a rise in the values of  $\epsilon$  and  $\tan\delta$ .  $\epsilon$  of ordinary silicate glasses and borate glasses lies between 6 and 10, of silicate glasses with high (50 mole%)  $\text{PbO}$  content between 17 and 18. G. I. Skanavi and A. M. Kashtanova (ZhFT, XXVII, 1770, 1957) obtained devitrified boron-lead-titanium glasses with  $\epsilon = 35$ . They explained the high  $\epsilon$  value as being due to the formation of crystalline lead titanate. Glasses on the basis of  $\text{TeO}_2$  have  $\epsilon$  values of 28-32.

Glasses on the basis of bismuth oxide have the highest  $\epsilon$  of all inorganic glasses known; their  $\tan\delta$  has the same order of magnitude as in alkali-free silicate glasses. Their use in industry appears very promising. There are

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B102/B214

Glasses with high dielectric constant

3 figures and 4 references: 1 Soviet-bloc and 3 non-Soviet-bloc. The three references to English-language publications read as follows: H. M. Heaton, H. Moore. J. Soc. Glass Techn. 41, 3, 1957; M. Heynes, H. Rowson, J. Soc. Glass Techn., 41, 347, 1957; J. Ph. Poley. Nature, 174, No. 4423, 268, 1954.

ASSOCIATION: Institut khimii silikatov AN SSSR Leningrad (Institute of Silicate Chemistry of the AS USSR, Leningrad)

SUBMITTED: February 24, 1961

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Glasses with high dielectric constant

S/181/61/003/006/031/031  
B102/B214

Legend to Fig. 1:  
 ▲ devitrificating and  
 crystallizing glasses  
 • transparent, light  
 yellow glasses.

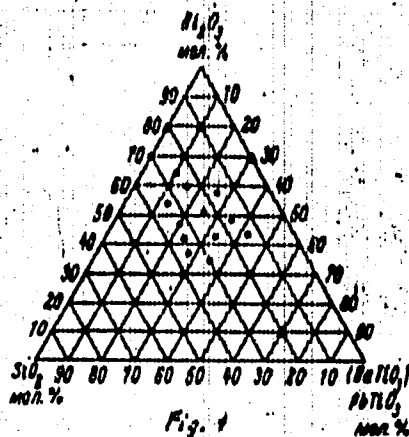


Fig. 1

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